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| 1 | BRS | L14 | 0 | HCPN-11 | USPAT; US-PGPUB; EPO; JPO; | 2004/01/02 10:02 | | | 0 |

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LE 'CAPLUS' ENTERED AT 10:05:09 ON 02 JAN 2004
E IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
EASE SEE "HELP USAGETERMS" FOR DETAILS.
PYRIGHT (C) 2004 AMERICAN CHEMICAL SOCIETY (ACS)
LE 'BIOSIS' ENTERED AT 10:05:09 ON 02 JAN 2004
PYRIGHT (C) 2004 BIOLOGICAL ABSTRACTS INC.(R)
LE 'EMBASE' ENTERED AT 10:05:09 ON 02 JAN 2004
PYRIGHT (C) 2004 Elsevier Inc. All rights reserved.
LE 'SCISEARCH' ENTERED AT 10:05:09 ON 02 JAN 2004
PYRIGHT 2004 THOMSON ISI
LE 'AGRICOLA' ENTERED AT 10:05:09 ON 02 JAN 2004
s human chaperone protein
              8 HUMAN CHAPERONE PROTEIN
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              0 HCPN-11
duplicate remove 11
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EP DUPLICATES FROM MORE THAN ONE FILE? Y/(N):n
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                             2001:101182 CAPLUS
CESSION NUMBER:
                             134:159191
CUMENT NUMBER:
                                                                              ***protein***
                                ***Human***
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TLE:
                             (HCPN) sequence homologs, their sequences, cDNA encoding them, and their biological and therapeutic
                             Yue, Henry; Bandman, Olga; Tang, Y. Tom; Baughn,
Mariah R.; Azimzai, Yalda; Lu, Dyung Aina M.
IVENTOR(S):
                             Incyte Genomics, Inc., USA
ATENT ASSIGNEE(S):
                             PCT Int. Appl., 101 pp.
OURCE:
                             CODEN: PIXXD2
                             Patent
CUMENT TYPE:
                             English
NGUAGE:
AMILY ACC. NUM. COUNT:
ATENT INFORMATION:
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115 A2 20020508 EP 2000-952500 20000803
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29327 T2 20031007 JP 2
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                                                                           19990803
RIORITY APPLN. INFO.:
                                                 US 1999-160924P
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    The invention provides eleven human proteins, which are believed to be
    chaperone proteins (HCPN) based on sequence homol. to known heat-shock,
    chaperone and DnaJ proteins. The invention also provides cDNA mols.
    encoding the HCPN sequence homologs. The invention further provides a DNA
    construct contg. a promoter linked to said HCPN cDNA mols., and a cell and/or organism transformed for with said DNA construct, which are used
    for recombinant prodn. of HCPN. Still further the invention provides: (1) a pharmaceutical compon conton said HCPN: (2) antibodies specific for
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HCPN; (3) primers and/or probes specific for polynucleotides encoding HCPN; (4) RNA equiv. of HCPN cDNA mols.; and (5) mol. genetic techniques, such as polymerase chain reaction (PCR) and/or nucleic acid hybridization for detecting polynucleotides encoding HCPN using said primers and probes. Finally the invention provides: (1) screening methods for agonists and/or antagonists of HCPN, and (2) use of identified agonists and/or antagonists in treating a disease or disorder assocd. With an imbalance of functional The CDNA sequences as well as the corresponding amino acid sequences of the HCPN sequence homologs are provided. The invention presented information on the cloning of each cDNA mol., including what tissues were utilized in constructing the cDNA libraries. In addn., the invention presented information on the structure and potential function of the HCPN including: (1) potential phosphorylation and glycosylation sites; (2) signature sequences and protein motifs; and (3) proteins from other organisms that show homol. Finally, the invention presented information on the tissue expression of the cDNA clones as detd. by Northern blot, and diseases and/or disorders assocd with these tissues

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diseases and/or disorders assocd. with these tissues.
   ANSWER 2 OF 4 CAPLUS COPYRIGHT 2004 ACS ON STN
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CESSION NUMBER:
CUMENT NUMBER:
                           132:247181
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                              ***Human***
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TLE:
                           and their encoding nucleic acids
                           Tang, Y. Tom; Hillman, Jennifer L.; Yue, Henry; Patterson, Chandra; Baughn, Mariah R.; Batra, Sajeev
IVENTOR(S):
                            Incyte Pharmaceuticals, Inc., USA
ATENT ASSIGNEE(S):
                           PCT Int. Appl., 88 pp.
OURCE:
                           CODEN: PIXXD2
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OCUMENT TYPE:
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ANGUAGE:
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US 1999-294698
WO 1999-US22027
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                                                                 W
                                    ***human***
    The invention provides 6
***proteins*** (HCHP)
                                                       ***chaperone***
                          (HCHP) and polynucleotides which identify and encode
    HCHP. The invention also provides expression vectors, host cells, antibodies, agonists, and antagonists. The invention also provides the
    use of these sequences in the diagnosis, treatment, and prevention of
    neurodegenerative, metabolic, developmental, autoimmune-inflammatory
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expression of HCHP. DUPLICATE 1 MEDLINE on STN ANSWER 3 OF 4 CCESSION NUMBER: 1999024006 MEDLINE OCUMENT NUMBER:

PubMed ID: 9804845 99024006

Human Hsp70 and Hsp40 chaperone proteins facilitate human papillomavirus-11 E1 protein binding to the origin and stimulate cell-free DNA replication.

disorders and cell proliferative disorders including cancer assocd. with

Liu J S; Kuo S R; Makhov A M; Cyr D M; Griffith J D; Broker

TR; Chow L T

ITLE:

UTHOR:

Department of Biochemistry and Molecular Genetics, ORPORATE SOURCE: University of Alabama at Birmingham, Birmingham, Alabama

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CA19014 (NCI)
NTRACT NUMBER:
   CA36200 (NCI)
   GM31819 (NIGMS)
                        JOURNAL OF BIOLOGICAL CHEMISTRY, (1998 Nov 13) 273 (46)
URCE:
                        30704-12.
                        Journal code: 2985121R. ISSN: 0021-9258.
                       United States
B. COUNTRY:
                        Journal; Article; (JOURNAL ARTICLE)
CUMENT TYPE:
                       English
NGUAGE:
                       Priority Journals
LE SEGMENT:
TRY MONTH:
                        199812
                        Entered STN: 19990115
ITRY DATE:
                       Last Updated on STN: 19990115
Entered Medline: 19981208
    Human papillomavirus replication initiator, the E1 helicase, binds weakly to the origin of DNA replication. Purified ***human***

***chaperone*** ***proteins*** Hsp70 and Hsp40 (HDJ-1 and HDJ-2)
    independently and additively enhanced E1 binding to the origin.
    interaction between E1 and Hsp70 was transient and required ATP
    hydrolysis, whereas Hsp40 bound to E1 directly and remained in the
    complex. A peptide of 20 residues spanning the HPD loop and helix II of
    the J domain of YDJ-1 also stimulated E1 binding to the origin, alone or
    in combination with Hsp70 or Hsp40. A mutated peptide (H34Q) had a reduced activity, while an adjacent or an overlapping peptide had no effect. Neither Hsp70 nor the J peptide altered the E1/DNA ratio in the complex. Electron microscopy showed that E1 mainly bound to DNA as a hexamer. In the presence of Hsp40, E1 primarily bound to DNA as a dihexamer. Preincubation of chaperones with viral E1 and template shortened the lag time and increased replication in a cell-free system.
    shortened the lag time and increased replication in a cell-free system.
    Since two helicases are essential for bidirectional replication of human
    papillomavirus DNA, these results demonstrate that, as in prokaryotes, chaperones play an important role in the assembly of preinitiation
    complexes on the origin.
    ANSWER 4 OF 4 SCISEARCH COPYRIGHT 2004 THOMSON ISI ON STN
CCESSION NUMBER:
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                                                                    ***CHAPERONE***
ITLE:
                            ***PROTEIN***
                                                 TO REPRODUCE
                         BROWN P (Reprint)
JTHOR:
                         NEW SCIENTIST, (03 DEC 1994) Vol. 144, No. 1954, pp. 20.
OURCE:
                         ISSN: 0262-4079.
OCUMENT TYPE:
                         Editorial; Journal
ILE SEGMENT:
                         AGRI; ENGI
                         ENGLISH
ANGUAGE:
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EFERENCE COUNT:
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CESSION NUMBER:
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CUMENT NUMBER:
                                                              Human chaperone protein (HCPN) sequence homologs,
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                                                              biological and therapeutic uses
                                                              Yue, Henry; Bandman, Olga; Tang, Y. Tom; Baughn,
Mariah R.; Azimzai, Yalda; Lu, Dyung Aina M.
VENTOR(S):
                                                              Incyte Genomics, Inc., USA
TENT ASSIGNEE(S):
                                                              PCT Int. Appl., 101 pp.
URCE:
                                                              CODEN: PIXXD2
CUMENT TYPE:
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NGUAGE:
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1203015

A2 20020508

EP 2000-952500

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        The invention provides eleven human proteins, which are believed to be chaperone proteins (HCPN) based on sequence homol. to known heat-shock, chaperone and DnaJ proteins. The invention also provides cDNA mols.
        encoding the HCPN sequence homologs. The invention further provides a DNA construct contg. a promoter linked to said HCPN cDNA mols., and a cell and/or organism transformed for with said DNA construct, which are used
         for recombinant prodn. of HCPN. Still further the invention provides: (1)
        Tor recombinant prodn. of HCPN. Still further the invention provides: (1) a pharmaceutical compn. contg. said HCPN; (2) antibodies specific for HCPN; (3) primers and/or probes specific for polynucleotides encoding HCPN; (4) RNA equiv. of HCPN cDNA mols.; and (5) mol. genetic techniques, such as polymerase chain reaction (PCR) and/or nucleic acid hybridization for detecting polynucleotides encoding HCPN using said primers and probes. Finally the invention provides: (1) screening methods for agonists and/or antagonists of HCPN, and (2) use of identified agonists and/or antagonists in treating a disease or disorder assocd. With an imbalance of functional HCPN. The cDNA sequences as well as the corresponding amino acid
                           The CDNA sequences as well as the corresponding amino acid
         sequences of the HCPN sequence homologs are provided. The invention
         presented information on the cloning of each cDNA mol., including what
          tissues were utilized in constructing the cDNA libraries. In addn., the
         invention presented information on the structure and potential function of
         the HCPN including: (1) potential phosphorylation and glycosylation sites; (2) signature sequences and protein motifs; and (3) proteins from other organisms that show homol. Finally, the invention presented information on the tissue expression of the cDNA clones as detd. by Northern blot, and dispasses and/or d
         diseases and/or disorders assocd. with these tissues.
> s yue henry/au
                          411 YUE HENRY/AU
> s bandman olga/au
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s lu dyung/au 0 LU DYUNG/AU s (16 or 17 or 18 or 19 or 110) 11546 (L6 OR L7 OR L8 OR L9 OR L10) > s 112 and 11 2 L12 AND L1 duplicate remove 113 ROCESSING COMPLETED FOR L13 2 DUPLICATE REMOVE L13 (0 DUPLICATES REMOVED) ⊳ d 114 1-2 ibib abs 14 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2004 ACS ON STN 2001:101182 CAPLUS CCESSION NUMBER: OCUMENT NUMBER: 134:159191 ***Human*** ***chaperone*** ***protein*** TLE: (HCPN) sequence homologs, their sequences, cDNA encoding them, and their biological and therapeutic ***Yue, Henry*** ; ***Bandman, Olga***
Tang, Y. Tom ; Baughn, Mariah R.;
Yalda*** ; Lu, Dyung Aina M. NVENTOR(S): ***Azimzai,*** Incyte Genomics, Inc., USA ATENT ASSIGNEE(S): PCT Int. Appl., 101 pp. OURCE: CODEN: PIXXD2 OCUMENT TYPE: Patent ANGUAGE: English AMILY ACC. NUM. COUNT: ATENT INFORMATION: KIND DATE APPLICATION NO. DATE PATENT NO. wo 2001009178 Α2 20010208 wo 2000-us21313 20000803 Α3 20010927 wo 2001009178 AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG EP 2000-952500 20000803 EP 1203015 AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL 329327 T2 20031007 JP 2001-513984 20000803 JP 2003529327 US 1999-146908P P RIORITY APPLN. INFO.: US 1999-160924P P 19991022 wo 2000-us21313 w 20000803 The invention provides eleven human proteins, which are believed to be chaperone proteins (HCPN) based on sequence homol to known heat-shock, chaperone and DnaJ proteins. The invention also provides cDNA mols. encoding the HCPN sequence homologs. The invention further provides a DNA construct contg. a promoter linked to said HCPN cDNA mols., and a cell and/or organism transformed for with said DNA construct, which are used for proceedings and the construct of the con for recombinant prodn. of HCPN. Still further the invention provides: (1) a pharmaceutical compn. contg. said HCPN; (2) antibodies specific for HCPN; (3) primers and/or probes specific for polynucleotides encoding HCPN; (4) RNA equiv. of HCPN cDNA mols.; and (5) mol. genetic techniques, such as polymerase chain reaction (PCR) and/or nucleic acid hybridization for detecting polynucleotides encoding HCPN using said primers and probes. Finally the invention provides: (1) screening methods for agonists and/or antagonists of HCPN, and (2) use of identified agonists and/or antagonists in treating a disease or disorder assocd. With an imbalance of functional HCPN. The cDNA sequences as well as the corresponding amino acid

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on the tissue expression of the CDNA clones as detd. by Northern blot, and
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  ANSWER 2 OF 2 CAPLUS COPYRIGHT 2004 ACS on STN
                           2000:210378 CAPLUS
CESSION NUMBER:
                           132:247181
CUMENT NUMBER:
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                                                 ***chaperone***
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TLE:
                           and their encoding nucleic acids
                              ***Tang, Y. Tom*** ; Hillman, Jennifer L.;
VENTOR(S):
                              ***Yue, Henry*** ; Patterson, Chandra; Baughn, Mariah
                           R.; Batra, Sajeev
                           Incyte Pharmaceuticals, Inc., USA
TENT ASSIGNEE(S):
                           PCT Int. Appl., 88 pp.
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                                                       ***chaperone***
    The invention provides 6
                                    ***human***
      ***proteins*** (HCHP) and polynucleotides which identify and encode
    HCHP. The invention also provides expression vectors, host cells, antibodies, agonists, and antagonists. The invention also provides the use of these sequences in the diagnosis, treatment, and prevention of
    neurodegenerative, metabolic, developmental, autoimmune-inflammatory disorders and cell proliferative disorders including cancer assocd. with
    expression of HCHP.
> d his
    (FILE 'HOME' ENTERED AT 10:04:39 ON 02 JAN 2004)
    FILE 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH, AGRICOLA' ENTERED AT 10:05:09 ON 02 JAN 2004
               8 S HUMAN CHAPERONE PROTEIN
               0 S HCPN-11
               4 DUPLICATE REMOVE L1 (4 DUPLICATES REMOVED)
               O S POLUNUCLEOTIDE (P) L3
               1 S CELL (P) TRANSFORM? (P) L3
             411 S YUE HENRY/AU
            356 S BANDMAN OLGA/AU
          11140 S TANG Y?/AU
              16 S BAUGHN MARIAH/AU
             146 S AZIMZAI YALDA/AU
               0 S LU DYUNG/AU
          11546 S (L6 OR L7 OR L8 OR L9 OR L10)
               2 S L12 AND L1
```

2 DUPLICATE REMOVE L13 (0 DUPLICATES REMOVED)

> log y OST IN U.S. DOLLARS

12

SINCE FILE TOTAL **SESSTON** FNTRY